NOV 21 2006

This report is required by law (7 USC 2143). Failure to report according to the regulations can result in an order to cease and desist and to be subject to penalties as provided for in Section 2150.

See reverse side for additional information.

Interagency Report Control No 0180-DOA-AN

UNITED STATES DEPARTMENT OF AGRICULTURE
ANIMAL AND PLANT HEALTH INSPECTION SERVICE

ANNUAL REPORT OF RESEARCH FACILITY

(TYPE OR PRINT)

1. REGISTRATION NO. 31-R-0117 **CUSTOMER NO.** 28786

FORM APPROVED
OMB NO. 0579-0036

2. HEADQUARTERS RESEARCH FACILITY *(Name and Address, as registered with USDA, include Zip Code)*

WIL RESEARCH LABORATORIES LLC
1407 GEORGE ROAD
ASHLAND, OH 44805
(202) 434-7481

3. REPORTING FACILITY (List all locations where animals were housed or used in actual research, testing, teaching, or experimentation, or held for these purposes. Attach additional sheets if necessary.)

FACILITY LOCATIONS*(sites)*

See Attached Listing

REPORT OF ANIMALS USED BY OR UNDER CONTROL OF RESEARCH FACILITY *(Attach additional sheets if necessary or use APHIS FORM 7023A)*

A. Animals Covered By The Animal Welfare Regulations	B. Number of animals being bred, conditioned, or held for use in teaching, testing, experiments, research, or surgery but not yet used for such purposes.	C. Number of animals upon which teaching, research, experiments, or tests were conducted involving no pain, distress, or use of pain-relieving drugs.	D. Number of animals upon which experiments, teaching, research, surgery, or tests were conducted involving accompanying pain or distress to the animals and for which appropriate anesthetic, analgesic, or tranquilizing drugs were used.	E. Number of animals upon which teaching, experiments, research, surgery or tests were conducted involving accompanying pain or distress to the animals and for which the use of appropriate anesthetic,analgesic, or tranquilizing drugs would have adversely affected the procedures, results, or interpretation of the teaching, research, experiments, surgery, or tests. *(An explanation of the procedures producing pain or distress in these animals and the reasons such drugs were not used must be attached to this report)*	F. TOTAL NO. OF ANIMALS (Cols. C + D + E)
4. Dogs	185	780	83	0	863
5. Cats	0	28	0	0	28
6. Guinea Pigs	0	600	0	2	602
7. Hamsters	0	116	2	0	118
8. Rabbits	554	2510	171	19	2700
9. Non-Human Primates	207	583	43	0	626
10. Sheep	0	2	15	0	17
11. Pigs	92	79	0	0	79
12. Other Farm Animals					
Goats	0	4	1	0	5
13. Other Animals					

ASSURANCE STATEMENTS

1) Professionally acceptable standards governing the care, treatment, and use of animals, including appropriate use of anesthetic, analgesic, and tranquilizing drugs, prior to, during, and following actual research, teaching, testing, surgery, or experimentation were followed by this research facility.

2) Each principal investigator has considered alternatives to painful procedures.

3) This facility is adhering to the standards and regulations under the Act, and it has required that exceptions to the standards and regulations be specified and explained by the principal investigator and approved by the Institutional Animal Care and Use Committee (IACUC). **A summary of all the exceptions is attached to this annual report.** In addition to identifying the IACUC-approved exceptions, this summary includes a brief explanation of the exceptions, as well as the species and number of animals affected.

4) The attending veterinarian for this research facility has appropriate authority to ensure the provision of adequate veterinary care and to oversee the adequacy of other aspects of animal care and use.

CERTIFICATION BY HEADQUARTERS RESEARCH FACILITY OFFICIAL
(Chief Executive Officer or Legally Responsible Institutional official)

I certify that the above is true, correct, and complete (7 U.S.C. Section 2143)

SIGNATURE O

(b)(6), (b)(7)c

DATE SIGNED 11/20/2006

APHIS FORM 7023
(AUG 91)

(Replaces VS FORM 18-23 (Oct 88), which is obsolete

PART 1 - HEADQUARTERS

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2006 Category E Animals

Two (2) Guinea Pigs were placed in Category 'E".

Two (2) Guinea Pigs may have experienced pain and/or distress during the conduct of a guinea pig dermal maximization study and served as the positive controls. These two guinea pigs were involved in a study that was designed and conducted in compliance with EPA OPPTS Guideline 870.2600 (2003) and OECD Guidelines for Testing Chemicals, section 406 (1992). Dermal observations were made daily. The guinea pigs were evaluated daily by animal health technicians and other signs of pain and/or distress such as inappetence, weight loss or vocalization were not present.

Nineteen (19) rabbits were placed in Category "E".

Sixteen (16) rabbits may have experienced pain and /or distress while involved in a total of seven acute ocular irritation studies. These studies were designed and conducted in compliance with EPA OPPTS Guideline 870.2400 (1998) and OECD guideline 405 (2002). The purpose of these studies was to evaluate the irritative effects of the test materials when applied topically to the eye as a single administration to rabbits. The rabbits were evaluated daily by animal health technicians and the treated eyes were graded for severity of conjunctivitis, ocular discharge, hemorrhage of conjunctival tissue and degree of corneal damage. None of these rabbits showed signs of inappetence, weight loss or vocalization.

Three (3) rabbits were involved in two different acute dermal irritation studies and may have suffered pain and/or distress from corrosive skin lesions. The purpose of this study was to evaluate the irritative effects of the test material when applied topically to the skin as a single administration to rabbits. This study was designed and conducted in compliance with EPA OPPTS guideline 870.2500 (1998) and OECD Guidelines for Testing Chemicals, section 404 (2002). These rabbits were evaluated daily by animal health technicians and the condition of the skin was graded for the severity of erythema, edema, eschar formation and fissuring present at the application site. Other signs of pain and/or distress such as inappetence, weight loss, or vocalization were not observed.

WIL Research Laboratories, LLC

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IACUC approved reportable exceptions to the Animal Welfare Act

70 Beagle dogs used in Telemetry Cardiovascular Assessment Studies were implanted with DSI telemetry implants and exempted from exercise during the 14 day surgical recovery period. These animals were given regular opportunity for exercise and socialization prior to and after a 14 day surgical recovery. During the surgical recovery period, dogs were given daily contact and interaction with the technical staff and well as visual, auditory and olfactory stimuli from other dogs in the room.